ALLERGY RELIEF- loratadine tablet FSA STORE INC.

1193A-CRM-2023-0111

Drug Facts

Active ingredient (in each tablet)

Loratadine 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding, ask a health professional before use

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of age	ask a doctor

consumers with liver or kidney disease ask a doctor

Other information

- store between 20 to 25°C (68 to 77°F)
- retain carton for complete product information and warnings

Inactive ingredients

lactose monohydrate, magnesium stearate, pregelatinized starch (maize), sodium starch glycolate

Questions or comments?

1-844-705-4384

PRINCIPAL DISPLAY PANEL

NDC 81522-393-11

caring mill™

†Compare to the active ingeedient of Claritin® Tablets

Allergy Relief

Loratadine Tablets, 10 mg

Antihistamine

25 Hour

Relief of:

- Sneezing Runny nose Itchy, watery eyes Itchy throat or nose
- Non-drowsy*
- Original prescription strength
- Indoor & outdoor allergies

90 TABLETS

Actual Size

*When taken as directed. See Drug Facts Panel



ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81522-393
Route of Administration	ORAL		

Active Ingredient/Active Moiety Ingredient Name Basis of Strength LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN) LORATADINE 10 mg

Inactive Ingredients			
Ingredient Name	Strength		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
STARCH, CORN (UNII: O8232NY3SJ)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			

Product Characteristics			
Color	white (white to off white)	Score	no score
Shape	ROUND	Size	6mm

Flavor	Imprint Code	G;10
Contains		

F	Packaging				
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:81522- 393-11	1 in 1 CARTON	01/11/2023		
1		90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210722	01/11/2023	

Labeler - FSA STORE INC. (049283340)

Revised: 1/2023 FSA STORE INC.